



Cite this: *Phys. Chem. Chem. Phys.*,
2016, **18**, 31505

Extensive H-atom abstraction from benzoate by OH-radicals at the air–water interface†

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Much is known about OH-radical chemistry in the gas-phase and bulk water. Important atmospheric and biological processes, however, involve little investigated OH-radical reactions at aqueous interfaces with hydrophobic media. Here, we report the online mass-specific identification of the products and intermediates generated on the surface of aqueous (H₂O, D₂O) benzoate-h5 and -d5 microjets by ~8 ns •OH(g) pulses in air at 1 atm. Isotopic labeling lets us unambiguously identify the phenylperoxyl radicals that ensue H-abstraction from the aromatic ring and establish a lower bound (>26%) to this process as it takes place in the interfacial water nanolayers probed by our experiments. The significant extent of H-abstraction vs. its negligible contribution both in the gas-phase and bulk water underscores the unique properties of the air–water interface as a reaction medium. The enhancement of H-atom abstraction in interfacial water is ascribed, in part, to the relative destabilization of a more polar transition state for OH-radical addition vs. H-abstraction due to incomplete hydration at the low water densities prevalent therein.

Received 28th September 2016,
Accepted 1st November 2016

DOI: 10.1039/c6cp06652f

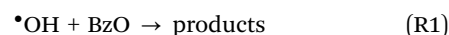
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Introduction

Benzoic acid (BA) is one of the most abundant carboxylic acid in the particulate matter (PM) found over most polluted urban areas. It has been recently reported that BA concentrations in PM_{2.5} collected over Beijing (average 1496 ng m⁻³ in Pekin University, 1278 ng m⁻³ in Yufa) exceed the concentrations of total diacids (~1010 ng m⁻³), fatty acids (~600 ng m⁻³) and ketocarboxylic acids (~120 ng m⁻³).¹ Low-volatility BA, which is produced both from direct traffic emissions and in the atmospheric oxidation of anthropogenic aromatic compounds, largely partitions to the aqueous phase where it reacts further with atmospheric oxidants.¹ BA (pK_a = 4.2) is largely present as benzoate (BzO) in atmospheric aqueous media. Since BzO is amphiphilic and relatively inert toward O₃ (*k* = 1.2 M⁻¹ s⁻¹),² the heterogeneous (interfacial) oxidation of BzO(aq) by •OH(g) is deemed to control its fate.^{3–8} Molecular dynamics (MD) calculations and surface-tension data confirm the affinity of BzO for aqueous surfaces.⁹ It has been recently realized that the photochemical aging of particulate organic matter is not only degradative but generates volatile organic compound (VOC) emissions and reactive species, such as hydroperoxides.^{10–12}

The identification of products and labile intermediates from the heterogeneous oxidation of organic matter in condensed phases has thus emerged as a major issue in the atmospheric chemistry of polluted urban air.^{6,13–15}

Here we address this issue and report direct, online mass-specific identification of the products of the oxidation of BzO(aq) by •OH(g) pulses on the surface of aqueous microjets (see Methods and Fig. S1, ESI†).¹¹ In such events, •OH(g) first sticks to the surface of water and is converted into hydrated •OH(H₂O)_{*n*} species,^{6,11,16} which react with BzO *via* (R1), or recombine into H₂O₂ (reaction (R9), Scheme 1) within interfacial layers:^{17,18}



Our technique can monitor *in situ* within ~1 ms the formation of primary products and intermediates on the surface of continuously flowing, uncontaminated aqueous surfaces at atmospheric pressure and 298 K.¹⁹

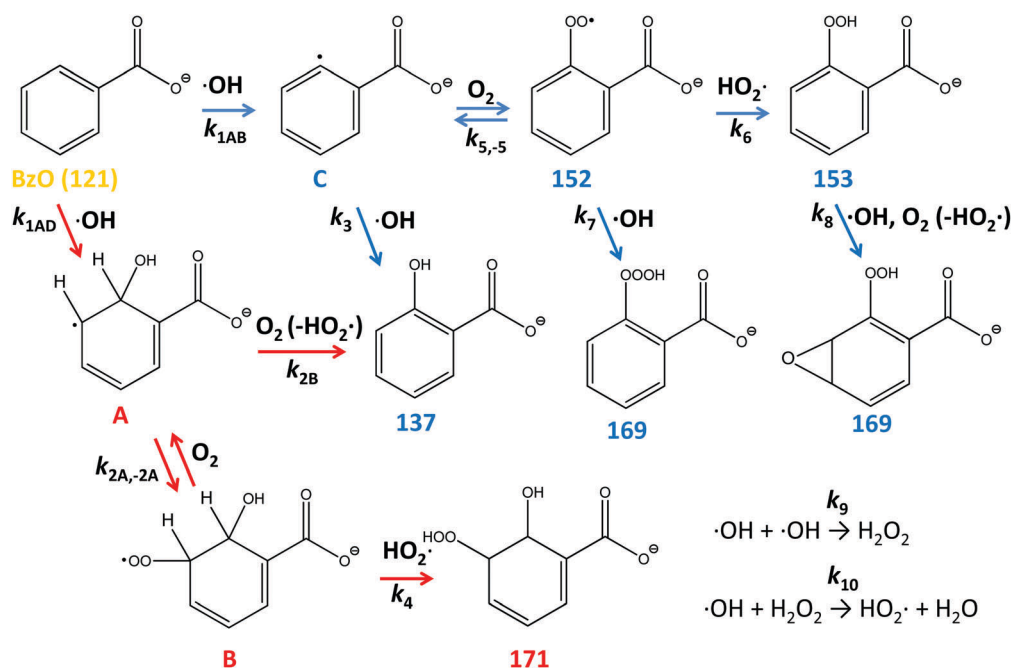
Experimental

The experimental setup has been described in previous publications.^{11,20} The prompt (within the ~10 μs lifetime of the intact microjets) formation of anionic products at the air–water interfaces of microjets from the reaction of aqueous reactants with gaseous OH-radicals at 1 atm at 298 K are monitored *in situ* by an electrospray ionization mass spectrometry (ES-MS, Agilent 6130 Quadrupole LC/MS Electrospray System, see Fig. S1, ESI†).¹¹ Aqueous solutions are pumped

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† Electronic supplementary information (ESI) available: Additional data and experimental details. See DOI: 10.1039/c6cp06652f



Scheme 1 Mechanism of formation and proposed structures (among the various positional and/or functional isomers in each case) of the species generated in the $\cdot\text{OH}$ -initiated oxidation of benzoate at the air–water interface.

(100 $\mu\text{L min}^{-1}$) into the spraying chamber of the mass spectrometer through a grounded stainless steel needle (100 μm bore) coaxial with a sheath issuing nebulizer $\text{N}_2(\text{g})$ at high gas velocity ($v_g \sim 160 \text{ m s}^{-1}$).²⁰ The surface specificity of our experiments had been previously demonstrated.^{20–26} The depth (or thickness) of the sampled interfacial layers can be controlled by varying the nebulizer gas velocity v_g , as evidenced by the fact that both ion signal intensities and relative anion surface affinities increase with higher gas velocities v_g and extrapolate to zero as $v_g \rightarrow 0$.²⁰ The ions detected by our mass spectrometer are ions that: (1) were already present or produced by chemical reactions in the interfacial layers of microjets (see previous publications for further details),^{20,23,27,28} (2) become incorporated into charged microdroplets produced during the stripping of interfacial layers by the nebulizing gas, and (3) finally ejected to the gas-phase and admitted into the mass spectrometer section *via* a polarized inlet port positively biased at 3.5 kV relative to ground.

The dissociation of $\text{O}_3(\text{g})$ by unfocused 266 nm laser pulses (laser beam diameter 10 mm, beam divergence ≤ 1.5 mrad, pulse duration ~ 8 ns) into $\text{O}(^1\text{D})$, followed by the reaction of $\text{O}(^1\text{D})$ with $\text{H}_2\text{O}(\text{g})$, in competition with its deactivation by $\text{N}_2(\text{g})$ and $\text{O}_2(\text{g})$ into $\text{O}(^3\text{P})$, yields $\cdot\text{OH}(\text{g})$ within ~ 6 ns. Order of magnitude $\cdot\text{OH}(\text{g})$ concentrations were estimated as described in previous publication.¹¹

Results and discussion

Fig. 1 shows a typical negative ion electrospray mass spectrum obtained from 1.0 mM $\text{BzO}(\text{aq})$ microjets under $\text{O}_3(\text{g})/\text{O}_2(\text{g})$

$\text{H}_2\text{O}(\text{g})/\text{N}_2(\text{g})$ mixtures as such or after being irradiated with 266 nm laser pulses.

At pH 4.0, $\sim 40\%$ BA ($\text{pK}_a = 4.2$) is dissociated into detectable benzoate $\text{C}_6\text{H}_5\text{--COO}^-$ (BzO) at $m/z = 121$ (m/z in Thompson units throughout). Recall that neutral species are transparent to mass spectrometry (see above). We verified that in the absence of light, $\text{O}_3(\text{g})$ does not generate new product signals (Fig. S2, ESI[†]), in line with the inertness of BzO toward O_3 ($k_{\text{BzO}+\text{O}_3} = 1.2 \text{ M}^{-1} \text{ s}^{-1}$ in bulk water).² Upon 266 nm pulse irradiation of the inflowing $\text{O}_3(\text{g})/\text{O}_2(\text{g})/\text{H}_2\text{O}(\text{g})/\text{N}_2(\text{g})$ mixtures, which generates $\cdot\text{OH}(\text{g})$ *in situ* within 8 ns, we observe the partial depletion of BzO and the simultaneous appearance of new signals, which we therefore ascribe to products of $\cdot\text{OH}$ reactions with BzO . The same products, albeit in different proportions, were observed over the $[\text{BzO}] = 0.01\text{--}10 \text{ mM}$ range. We confirmed that reactant depletion and product formation require both the participation of $\text{O}_3(\text{g})$ and actinic 266 nm photons (Fig. S2 and S3, ESI[†]), that is, the chemistry we observe is neither due to benzoate ozonation or benzoate photolysis, but involves reactions of gas-phase OH -radicals with interfacial benzoate.

For the experiments shown in Fig. 1, we estimate $[\cdot\text{OH}(\text{g})]_0 \sim 2 \times 10^{14} \text{ molecules cm}^{-3}$ at 40 mJ pulse⁻¹ (the highest laser pulse energy used in our experiments) within the laser-irradiated volume.¹¹ $[\cdot\text{OH}(\text{g})]$ on the surface of microjets $\leq 1 \text{ mm}$ apart are estimated to be a factor of 10 lower. It might be argued that they are larger than atmospheric concentrations, but we note that reactant conversions during the $\tau \sim 10 \mu\text{s}$ reaction times of our experiments are comparable to those in which aerosol droplets exposed to typical tropospheric $[\cdot\text{OH}] \approx 10^6 \text{ molecules cm}^{-3}$ concentrations for 40 min. Product distributions, however, are expected to be different. In our experiments, yields of secondary

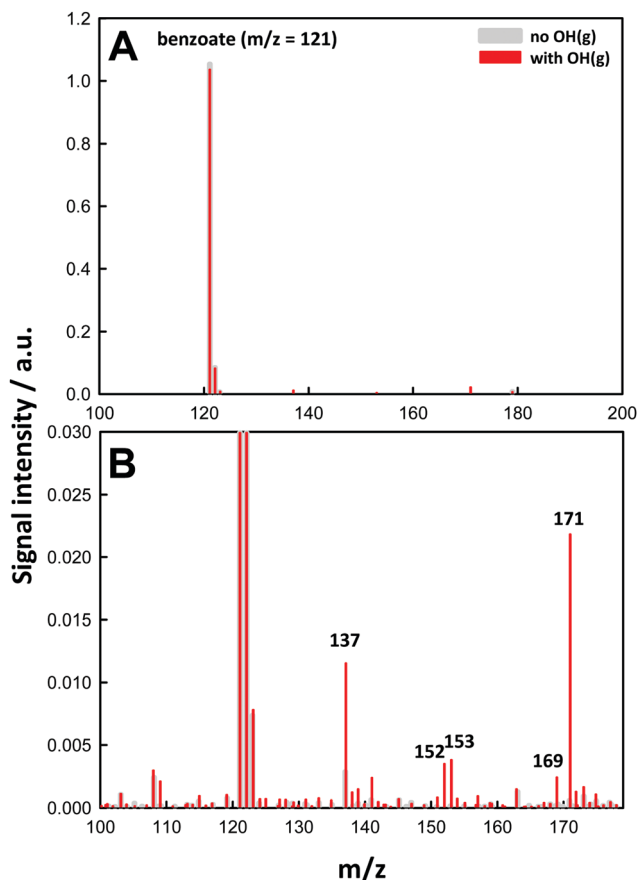


Fig. 1 (A) Negative ion electrospray mass spectra of 1.0 mM (pH 4.0) benzoic acid microjets in the presence of 2×10^{15} molecules cm^{-3} $\text{O}_3(\text{g})$ in $\text{O}_2(\text{g})/\text{H}_2\text{O}(\text{g})/\text{N}_2(\text{g})$ mixtures with the 266 nm laser pulses (40 mJ, ~ 8 ns, 10 Hz) off (gray) or on (red). (B) Zooming in the $\bullet\text{OH}$ -oxidation products. See text for details.

products resulting from organic radical + OH-radical reactions will be enhanced relative to those produced under atmospheric conditions.

The molecular formulas of the species generated in our experiments could be inferred from their mass-to-charge ratios. Thus, the $m/z = 137$ signal in Fig. 1B is readily assigned to hydroxy-benzoates $\text{C}_6\text{H}_4(\text{OH})\text{COO}^-$ (BzO-OH): $137 = 121 + 16 = \text{BzO} - \text{H} + \bullet\text{OH}$. It is important to note that BzO-OH can be produced *via* two reaction channels: one initiated by $\bullet\text{OH}$ -addition to BzO : $137 = 121 (\text{BzO}) + 17 (\bullet\text{OH}) + 32 (\text{O}_2) - 33 (\text{HO}_2\bullet)$, and another one initiated by H-abstraction from BzO : $137 = 121 (\text{BzO}) - 1 (\text{H}) + 17 (\bullet\text{OH})$ (Scheme 1). Note that one phenylic H-atom is removed in both cases. Similarly, the $m/z = 153$ signal corresponds to structures formally derived from O-addition to BzO-OH : $153 = 137 + 16$, such as benzoate hydroperoxides $\text{C}_6\text{H}_4(\text{OOH})\text{COO}^-$ (BzO-OOH) or di-hydroxy-benzoates $\text{C}_6\text{H}_3(\text{OH})_2\text{COO}^-$ ($\text{BzO}(\text{OH})_2$). The difference is that the hydroperoxides will retain 4 phenylic H-atoms but the di-hydroxy species only 3. The $m/z = 169$ signal corresponds to species formally derived from the addition of a third O-atom to BzO-OOH : $169 = 153 + 16$.^{29,30} The $m/z = 171$ signal corresponds to the only species produced by successive $\bullet\text{OH}$ and $\text{HO}_2\bullet$ additions: $171 = 121 (\text{BzO}) + 17 (\bullet\text{OH}) + 33$

($\text{HO}_2\bullet$), *via* processes that retain all 5 phenylic H-atoms. In addition to the above closed-shell species, we detected even mass $m/z = 152$ signals, which correspond to isomeric peroxy radicals $\text{C}_6\text{H}_4(\text{OO}\bullet)\text{COO}^-$ ($\text{BzO-O}_2\bullet$) derived from O_2 -addition to the phenylic radicals generated by H-abstraction from BzO : $152 = 121 (\text{BzO}) - 1 (\text{H}) + 32 (\text{O}_2)$.³¹ To our knowledge, this is the first report on the detection of early, labile intermediates, such as peroxy radicals and hydroperoxides, in the oxidation of aromatics by $\bullet\text{OH}$ at the air-water interface. Since mass spectrometry reports molecular mass, the structures shown in Scheme 1 stand for all possible positional isomers in each case.^{32–34}

Experiments in D_2O provided additional evidence on molecular assignments (Fig. 2). Thus, hydroxy-benzoates and hydroxy-hydroperoxides, which possess one and two exchangeable (O-)H-atoms, generate ($M + 1$) and ($M + 2$) species, respectively. We note that the H-containing nascent phenol and hydroperoxide groups are able to exchange with D_2O prior to detection ~ 1 ms later.^{10–12} The finding that $m/z = 152$ does not shift in D_2O solvent is clearly consistent with $\text{BzO-O}_2\bullet$ peroxy radical structural isomers.

Our assignments were validated in experiments involving isotopically labeled BzO-d5 (benzoate-2, 3, 4, 5, 6-d5) in H_2O and D_2O as solvents. Fig. 3A shows the mass spectra for the $\bullet\text{OH}$ oxidation of BzO-d5 ($m/z = 126$) in H_2O . In this case, we detected products at $m/z = 141, 156, 157, 173$ and 176 , which are consistent with the structures proposed in Scheme 1. For example, the hydroxy-benzoates signal shifts from $m/z = 137$ in BzO-h5 to 141 in BzO-d5 , *i.e.*, as expected from species derived from the abstraction of one D-atom from BzO-d5 . Similarly, D-abstraction precedes the formation of the benzoate peroxy radicals ($\text{BzO-O}_2\bullet$) and benzoate hydroperoxides (BzO-OOH), whose signals, originally at $m/z = 152$ and 153 , shift to 156 and 157 , respectively. The only species retaining all 5 D-atoms are those corresponding to hydroxy-hydroperoxides (Scheme 1), which shift from $m/z = 171$ to 176 as proof that they result from

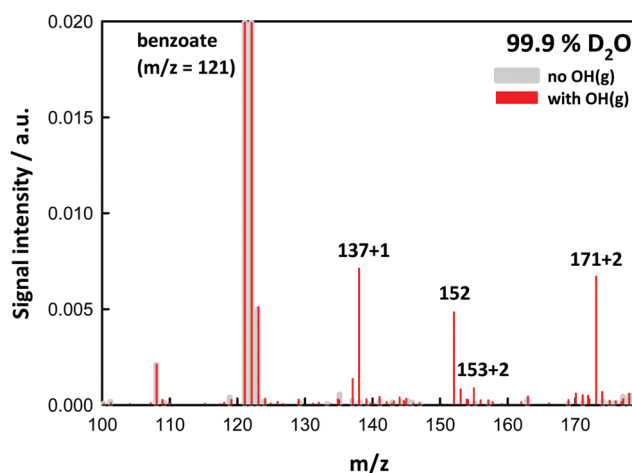


Fig. 2 Negative ion electrospray mass spectra of 0.5 mM benzoic acid in D_2O (99.9 atom % D) microjets exposed to 6.9×10^{15} molecules cm^{-3} $\text{O}_3(\text{g})$ in $\text{O}_2(\text{g})/\text{H}_2\text{O}(\text{g})/\text{N}_2(\text{g})$ mixtures at 1 atm and 298 K. Gray: laser off. Red: under 40 mJ, ~ 8 ns pulses (at 10 Hz) of 266 nm radiation.

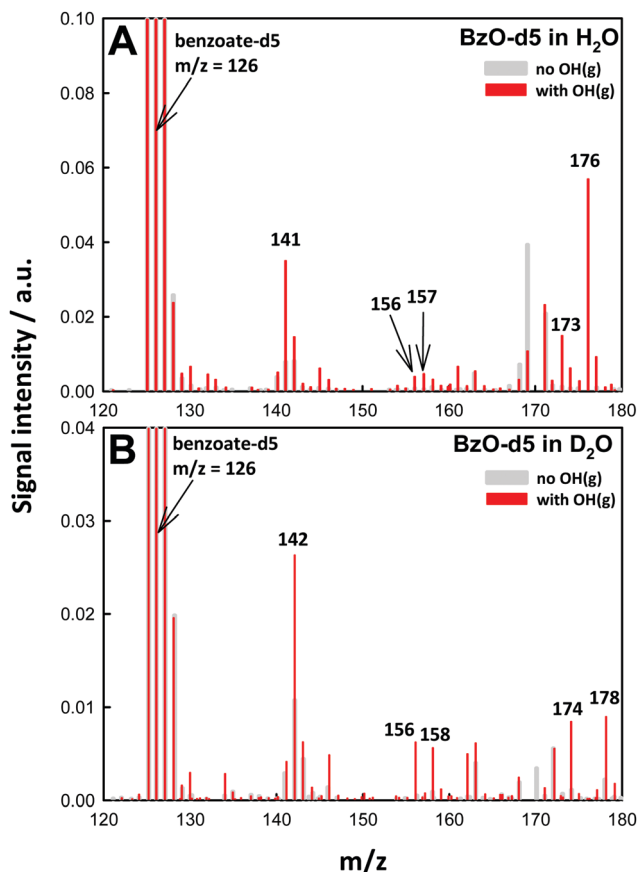


Fig. 3 (A) Negative ion electrospray mass spectra (background subtracted) of 10 mM C_6D_5COOH benzoic acid- d_5 microjets in H_2O . (B) in D_2O , in the presence of 2×10^{16} molecules cm^{-3} $O_3(g)$ at 1 atm and 298 K. Gray: laser off. Red: under 14 mJ, ~ 8 ns (at 10 Hz) 266 nm pulses.

OH-addition to the ring. The shift of the $m/z = 169$ signal in $BzO-h_5$ to $m/z = 173$ in $BzO-d_5$ excludes a tri-hydroxy-benzoate ($BzO-(OH)_3$) (which would have led to a $m/z = 169$ to 171 shift) but is consistent with trioxides or epoxide hydroperoxides (Scheme 1).

The $\bullet OH$ oxidation of $BzO-d_5$ in D_2O gives rise to $m/z = 142$, 156, 158, 174 and 178 products (Fig. 3B). The shift of the $m/z = 141$ signal in H_2O (Fig. 3A) to $m/z = 142$ in D_2O (Fig. 3B) is consistent with hydroxy-benzoates containing the exchangeable (O-)H-atom brought by $\bullet OH$. Also, according to the proposed structures, the signals assigned to the hydroxy-hydroperoxides, which contain two (O-)H-atoms, shift by two mass units from $m/z = 176$ to 178. As expected, signals assigned to benzoate peroxy radicals ($m/z = 156$ in Fig. 3A and B) do not shift as D_2O replaces H_2O as solvent. The $m/z = 173$ signals in Fig. 3A shift to $m/z = 174$ in D_2O , as expected from the presence of only one exchangeable H-atom in the proposed structures (Scheme 1) rather than to 176 should they correspond to tri-hydroxy isomers ($BzO-(OH)_3$). A benzoate trioxide ($m/z = 169$) could be formed by the radical-radical reaction ($BzO-O_2\bullet + \bullet OH$). The known negative temperature dependence of the ($CH_3O_2\bullet + \bullet OH = CH_3O\bullet + HO_2\bullet$) reaction in the gas-phase³⁵ is indicative of an associative reaction proceeding *via* a chemically activated

trioxide $CH_3OOOH\bullet$ intermediate. Thus, it is conceivable that a similar $BzO-OOOH\bullet$ trioxide could be formed and stabilized from ($BzO-O_2\bullet + \bullet OH$) in aqueous media. Alternatively, hydroperoxy epoxides (also $m/z = 169$) could be formed from the reaction of a hydroperoxide with $\bullet OH$, followed by O_2 addition and elimination of $HO_2\bullet$ (see Scheme 1). The formation of epoxides has been previously proposed in reactions of aromatics with $\bullet OH$ in the presence of O_2 .²⁹

We consider that our key finding is the detection of substantial yields of $BzO-O_2\bullet$ radicals, which quantify the extent of H-abstraction from the aromatic ring at the air-water interface. Our result is in marked contrast with previous studies in bulk water, in which addition was the exclusive channel in OH-radical reactions with aromatics.^{31,33,34} Our detection of labile intermediates and products, such as $BzO-O_2\bullet$ and the hydroperoxides *vis-à-vis* reports on the exclusive formation of hydroxybenzoates ($BzO-OH$) reported in the oxidation of BzO by $\bullet OH$ in bulk water,^{33,34} therefore implies that (1) all intermediates we detect within ~ 1 ms are ultimately converted to hydroxybenzoates, (2) they may have been missed in the other studies, or (3) the reaction proceeds by different mechanisms in bulk water *vs.* at the air-water interface. It should be emphasized that, at variance with the oxidation of alkyl-carboxylic acids by $\bullet OH$ under similar conditions,^{10–12} we found no evidence of putative products of $BzO-O_2\bullet$ self-reactions, such as alkoxy radicals and alcohol/carbonyls. Thus, self-reactions of the bulky secondary peroxy radicals $BzO-O_2\bullet$ are relatively slow in the timeframe of our experiments.³⁶

The dependences of signal intensities as functions of laser energy and benzoate concentration provide valuable mechanistic clues. Fig. 4 shows mass signals as functions of laser energy per pulse. We assume that $\bullet OH$ doses increase linearly with pulse energy at low energies before plateauing at high pulse energies. Note that these are not kinetic plots, *i.e.*, mass signals as functions of time. The fact that all species, *i.e.*, primary and secondary, appear at the lowest laser pulse energies, *i.e.*, at the lowest OH-radical concentrations, means that all reactions are very fast (and therefore nearly independent of temperature) and not limited by reactants under present conditions.

Interestingly, we note that the main products in the more dilute 0.5 mM BzO solutions are the $m/z = 171$ hydroxy-hydroperoxides (Fig. 4C), which, as we have seen, provide a true measure of the addition channel, whereas in 10 mM BzO the main products are the $m/z = 137$ hydroxybenzoates (Fig. 4F). Rough estimates OH-radical concentrations help to rationalize these findings. We estimate that a 40 mJ pulse generates $[OH(g)]_0 \approx 3 \times 10^{14}$ molecules cm^{-3} , which translate (on the basis of the kinetic theory of gases³⁷ and a mean $\bullet OH$ speed $c = 6.4 \times 10^4$ cm s^{-1} at 298 K) into $\sim 5 \times 10^{18}$ molecules cm^{-2} s^{-1} flux on the surface of the microjets.³⁸ From the reported thermal accommodation coefficient of $\bullet OH$ on water: $S \sim 0.95$,³⁹ we estimate that the maximum dose of $\bullet OH$ incorporated into the surface of aqueous microjets during ~ 8 ns pulses is: $N_{OH} \leq 4 \times 10^{10}$ radicals cm^{-2} .

The remarkable fact that BzO signals decay by $\sim 7\%$ in 0.5 mM solutions, and by $\sim 14\%$ in the 20 times more concentrated

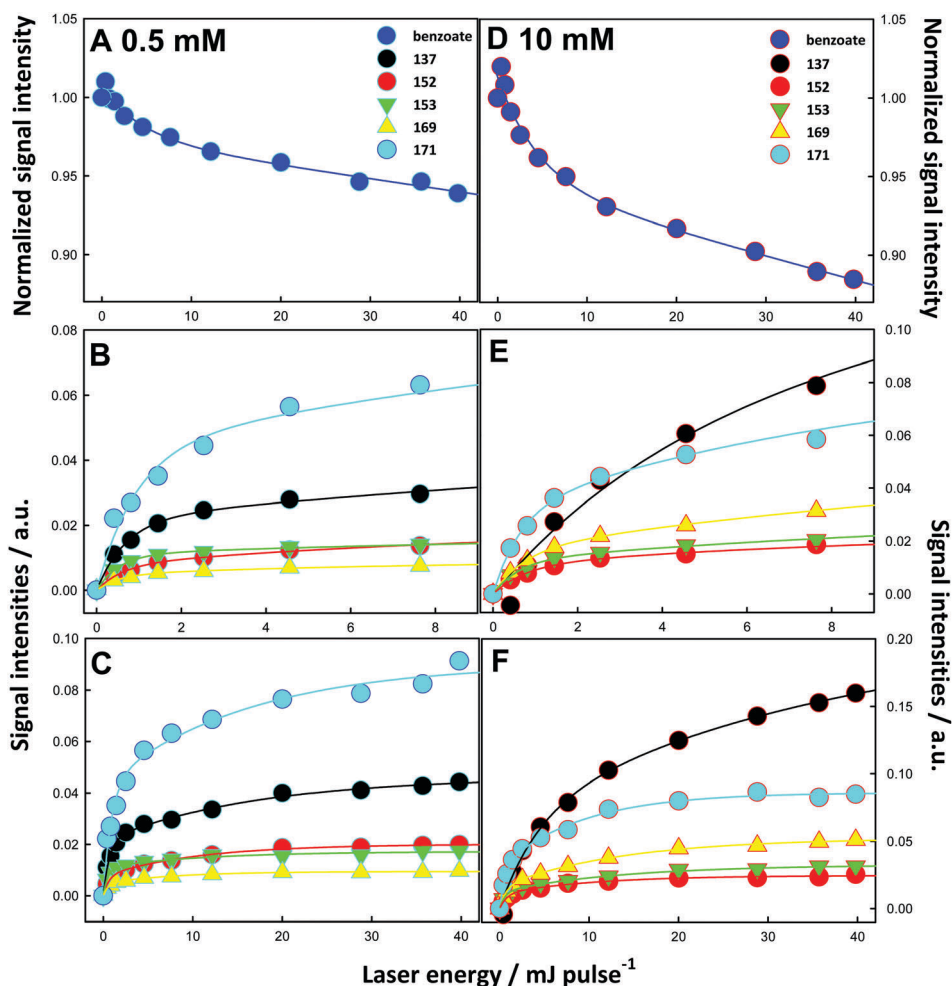


Fig. 4 (A) Reactant, (B and C) intermediates and products mass spectral signal intensities from aqueous 0.5 mM benzoic acid at pH 4.2 microjets, (D–F) from 10 mM benzoic acid at pH 3.4 microjets exposed to $O_3(g)/O_2(g)/H_2O(g)/N_2(g)$ mixtures, $[O_3(g)] \sim 4 \times 10^{15}$ molecules cm^{-3} , as functions of 266 nm laser energy (in $mJ\ pulse^{-1}$). Connecting lines are guides to the eye. See text for details.

10 mM solutions at the same $\bullet OH$ doses (Fig. 4A and D) has mechanistic implications. Also note that BzO signals do not decay as single exponentials but bottom out at large pulse energies (*i.e.*, at large $\bullet OH$ doses).^{10–12,40} The rate coefficients of ($\bullet OH + BzO$) *via* ($R_{1AB} + R_{1AD}$) and ($\bullet OH + \bullet OH$) *via* R_9 (Scheme 1) in bulk water are within a factor of two and correspond to diffusionally controlled reactions: $k_{1AB} + k_{1AD} = 2.5 \times 10^9\ M^{-1}\ s^{-1}$, $k_9 = 5.5 \times 10^9\ M^{-1}\ s^{-1}$.^{3,41,42} Since they are also expected to be fast and commensurate at the air–water interface, the above observations imply that: (1) $\bullet OH$ recombination into relatively inert H_2O_2 (*via* R_9 , Scheme 1) is competitive and more extensive than its reaction with BzO in the more dilute 0.5 mM BzO solutions, and (2) BzO must diffuse back from the bulk solution to replenish the depleted outermost layers. The first observation in turn requires that $\bullet OH$ and BzO should have comparable concentrations in the layers where these processes take place. This condition defines the average thickness δ of such layers. At the maximum dose of OH-radicals: $N_{\bullet OH} = 4 \times 10^{10}$ radicals cm^{-2} , the resulting $[\bullet OH]$ in layers of thickness δ is given by: $[\bullet OH] = N_{\bullet OH} \leq 4 \times 10^{10}$ radicals cm^{-2}/δ . By equating $[\bullet OH]$ to $[BzO] = 3 \times 10^{17}$ molecules cm^{-3}

(in 0.5 mM solutions), we derive a $\delta \leq 1.3$ nm value. It is apparent that our experiments effectively probe reactive events occurring in interfacial nanolayers.

An approximate but realistic estimate of the rates of the competing processes involved in BzO depletion reveals that reactions $R_{1AB} + R_{1AD}$ and R_9 take place in $\leq 1\ \mu s$ (see Kinetic Model Calculations in ESI† and Fig. S4 and S5). By assuming a typical value of diffusion coefficients in water: $D_{BzO} = 2 \times 10^{-5}\ cm^2\ s^{-1}$, we infer that BzO diffusion from the bulk into depleted layers takes place after chemical reactions are over. In fact, the slower time scale associated with replenishing BzO-depleted layers *via* diffusion sets an upper limit of $\sim 10\ \mu s$ to the lifetimes of intact microjets (*i.e.*, before they break into charged microdroplets). Longer lifetimes would have led to negligible BzO conversions because diffusion would have fully refilled interfacial layers. Shorter lifetimes, in contrast, would have preempted diffusion and led to exponential decays of BzO as a function of pulse energy. It should be emphasized that the much longer microjet lifetimes: $\sim 10\ \mu s$ *vs.* the ~ 8 ns laser pulses (which are shot every 100 ms) imply that our experiments

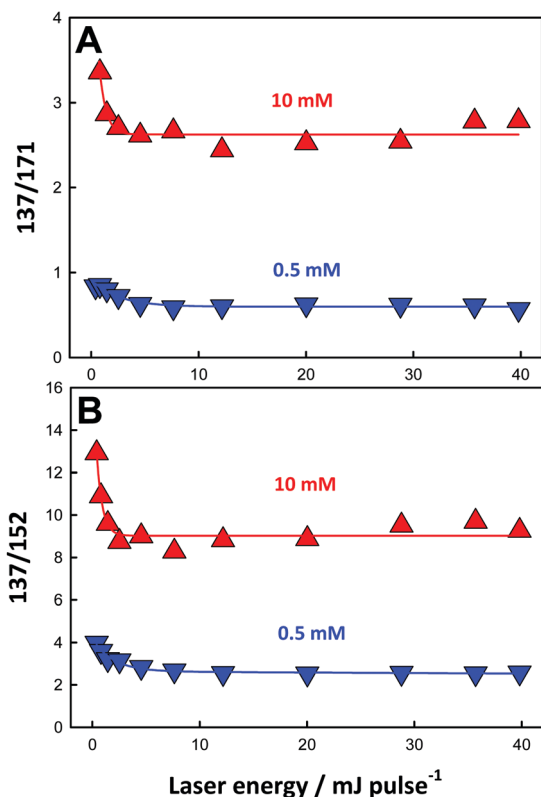


Fig. 5 (A) The 137/171 and (B) 137/152 ratios of signal intensities from aqueous 0.5 mM (blue triangles) and 10 mM (red triangles) benzoic acid microjets exposed to $\text{O}_3(\text{g})/\text{O}_2(\text{g})/\text{H}_2\text{O}(\text{g})/\text{N}_2(\text{g})$ mixtures as functions of 266 nm laser energy (in mJ pulse^{-1}). Connecting lines are guides to the eye. See text for details.

correspond to processes taking place in fresh air–water interfaces after being exposed once to strong, short $\bullet\text{OH}$ pulses.

The 137/171 and 137/152 ratios of signal intensities as a function of laser energy, *i.e.* $\bullet\text{OH}$ dose, are shown in Fig. 5A and B. The dependences of product yields, defined as the ratio of the individual signal intensities to the sum of product signal intensities, and the ratios of 137/171 and $(152 + 153 + 169)/(137 + 171)$ as functions of BzO concentration in experiments under 40 mJ pulses are shown in Fig. 6 and 7, respectively.

These trends are consistent with the mechanism outlined in Scheme 1. It should be realized that: (1) $m/z = 137$ species can be produced both *via* OH-radical addition and abstraction reactions in the sequences (1AD + 2B) and (1AB + 3), (2) whereas the $m/z = 171$ species can only be produced from OH-radical addition along the sequence (1AD + 2A + 4), *via* the hydroxycyclohexadienyls **A** (undetected) from reaction 1AD in equilibrium with peroxy radicals **B** (undetected).^{43,44} Since little H_2O_2 , the precursor of $\text{HO}_2\bullet$, will be produced at high [BzO] and low $\bullet\text{OH}$ doses, the low $[\text{HO}_2\bullet]/[\text{O}_2]$ ratios prevailing under such conditions will largely convert the addition intermediate **A** into the $m/z = 137$ hydroxy-benzoates rather than into the $m/z = 171$ hydroxy-hydroperoxide (Scheme 1). Note that neutral $\text{HO}_2\bullet$ is MS-silent. It is apparent that under such conditions the $m/z = 137$ hydroxybenzoates will be favored over the $m/z = 171$ hydroxy-hydroperoxide. Thus, the competition between $\bullet\text{OH}$

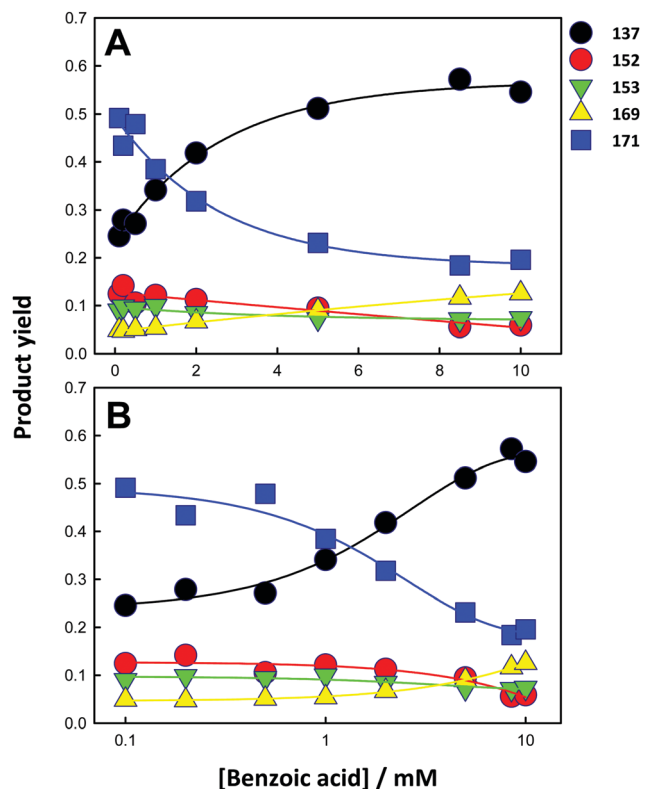


Fig. 6 (A) Fractional contributions of individual signal intensities to the mass spectra as functions of benzoic acid concentration in experiments under 40 mJ pulse^{-1} . (B) Semi-log plots. Connecting lines are guides to the eye.

reactions with BzO and recombination accounts for the higher 137/171 and 137/152 ratios observed at higher [BzO] and lower $\bullet\text{OH}$ doses (Fig. 5A and B), as well as the significant increase of the 137/171 ratio at larger [BzO] in Fig. 7.

On the basis of the preceding consideration, we estimate a lower limit to the branching between H-abstraction and addition from the ratio $\text{Abs/Add} \geq (152 + 153 + 169)/(137 + 171) = 0.35$. This is a lower limit to the extent of abstraction because the numerator is the sum of the signal intensities of the products derived from H-abstraction over those arising from addition by assuming that all 137 ensues from addition. It is apparent that at least 26% $[0.35/(1 + 0.35) = 0.26]$ of the $\bullet\text{OH}$ reacting with BzO will H-abtract from the aromatic ring at the air–water interface, independent of [BzO]. The ratio $\text{Abs/Add} \geq 0.35$ is consistent with a difference of $\Delta = E_{\text{Abs}} - E_{\text{Add}} \leq 0.6 \text{ kcal mol}^{-1}$ (from $\exp(-\Delta/RT) \geq 0.35$) between the activation energies for abstraction and addition at the air–water interface. We note that our $\Delta \leq 0.6 \text{ kcal mol}^{-1}$ value is significantly smaller than the $\Delta = 3 \text{ kcal mol}^{-1}$ reported for such competition in the presence of 2 water molecules,⁴⁵ and much smaller than the $\Delta = 6.5 \text{ kcal mol}^{-1}$ in the gas-phase, where H-abstraction is negligible.⁴⁶

The sensitivity of OH-radical reactions with aromatics to H-bonding with hydrophilic solvents is well established. An experimental and computational study of $\bullet\text{OH}$ reactions with aromatics in water and in polar, non-hydrophilic acetonitrile,⁴⁵ had reported rate coefficients ~ 65 times larger in water than in

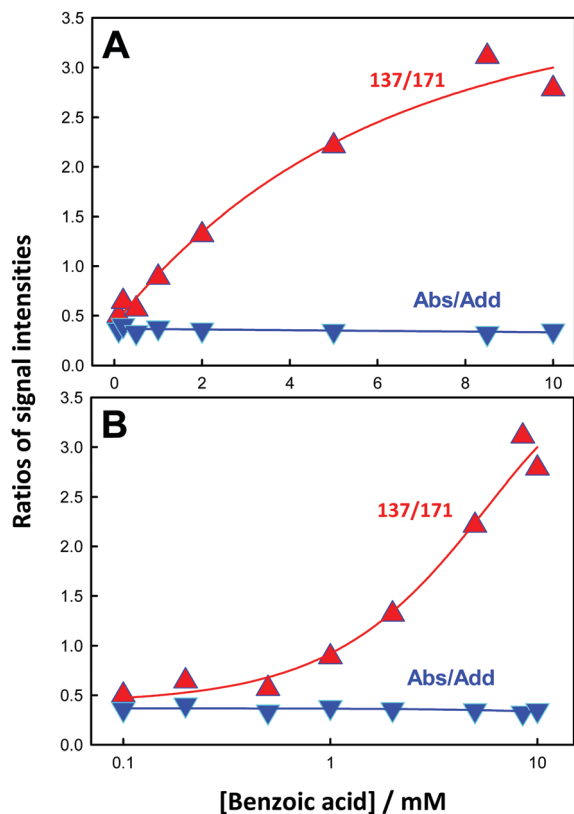


Fig. 7 (A) The 137/171 (red triangles) and $(152 + 153 + 169)/(137 + 171)$ (blue triangles) product signal ratios as functions of benzoic acid concentration in experiments under 40 mJ pulse^{-1} . (B) Semi-log plots. Connecting lines are guides to the eye. See text for details.

acetonitrile, as a clear indication of both the electrophilicity of the OH-radical and the polar nature of the transition states involved. Calculations also showed that the effects of binding one and two water molecules to reactants and transition states on free energies of activation were sensitive to relative orientation.^{47–50} In view of such information, we propose that H-abstraction is enhanced over addition in the anisotropic hydration environment provided by the air–water interface, where water density drops precipitously over 3 \AA .^{47–52} Molecular dynamics calculations based on many-body potentials that reliably simulate interfacial phenomena could shed further light on these processes.^{51,52}

The atmospheric implications of present findings are briefly discussed below. Our experiments simulate the heterogeneous oxidation of organic aerosol matter by gas-phase OH-radicals that stick to the surface to subsequently react at nearly diffusion controlled rates with aromatics therein. Significantly, they reveal that the oxidation of aqueous benzoate not only leads to hydroxylation but, *via* a sizable contribution of H-abstraction, to the formation of more reactive species, such as peroxy radicals and hydroperoxides, that can propagate radical chemistry *via* solar photolysis,^{53,54} or metal-catalyzed decomposition^{55,56} in the condensed aerosol phase. Given the hydrophobic character of benzoate, the most abundant organic species found in the $\text{PM}_{2.5}$ in polluted urban areas,¹ our experiments suggest that its oxidation by $\bullet\text{OH(g)}$ at the air–water interface may be an

important process in the photochemical aging of secondary organic aerosols.

Conclusion

Summing up, we report the first direct detection of peroxy radicals in significant yields during the oxidation of benzoate by OH-radicals at the air–water interface. They originate from significant ($>26\%$) H-abstraction from the aromatic ring, a pathway deemed to be absent in OH-radical reactions with aromatics in water. We also detected hydroperoxides and other hitherto unidentified products in addition to hydroxybenzoates. The significant extent of H-abstraction from the aromatic ring is tentatively ascribed to the relative destabilization of the more polar transition state for the OH-radical addition channel at the low water density prevalent in the outermost interfacial layers.

Acknowledgements

S. E. is grateful to the Hakubi Project of Kyoto University, the Iwatani Naoji Foundation's Research Grant and JSPS KAKENHI grant number 15H05328. M. R. H. and A. J. C. acknowledge support from the National Science Foundation (U.S.A.) Grant AC-1238977.

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